

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
17 May 2001 (17.05.2001)

PCT

(10) International Publication Number
WO 01/33975 A1

(51) International Patent Classification⁷: A23L 1/054,
1/0528, 1/0526, A61K 31/715, A61P 3/02

[BE/NL]; Kamperfoeliestraat 11, NL-6666 WS Heteren
(NL). BIJLSMA, Pieter, Brandt [NL/NL]; Elisabeth
Wolffstraat 65 e, NL-1043 TS Amsterdam (NL).

(21) International Application Number: PCT/NL00/00697

(74) Agent: JORRITSMA, Ruurd; Nederlandsch Octrooi-
bureau, Scheveningseweg 82, P.O. Box 29720, NL-2502 LS
The Hague (NL).

(22) International Filing Date:
29 September 2000 (29.09.2000)

(25) Filing Language: Dutch

(26) Publication Language: English

(30) Priority Data:
1013175 29 September 1999 (29.09.1999) NL

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ,
DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(71) Applicant (*for all designated States except US*): N.V. NU-
TRICIA [NL/NL]; P.O. Box 1, NL-2700 MA Zoetermeer
(NL).

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,
CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): KILIAAN,
Amanda, Johanne [NL/NL]; Harnjesweg 89, NL-6706
AS Wageningen (NL). GROOT, Jacques, Alphons
[NL/NL]; Heerenweg 153, NL-1851 KN Heiloo (NL).
TIMMERMANS, Johannes, Wilhelmus [NL/NL];
Schoonenburg 188, NL-6714 GE Ede (NL). VAN DER
MEULEN, Jan [NL/NL]; De Lipizzaner 33, NL-8252 GS
Dronten (NL). VAN LAERE, Katrien, Maria, Jozefa

Published:

— With international search report.

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: NUTRITIONAL COMPOSITIONS WHICH CONTAIN NON-DIGESTIBLE POLYSACCHARIDES AND USE
THEREOF TO REDUCE TRANSPORT THROUGH TIGHT JUNCTIONS

(57) Abstract: The present invention relates to the use of one or more non-digestible polysaccharides selected from the group con-
sisting of dextrans having a molecular weight of 8 kD to 40,000 kD, hydrolysed (gluco)mannans having a molecular weight of 0.5 kD
to 1,000 kD and hydrolysed (galacto)mannans having a molecular weight of 0.5 kD to 1,000 kD for the preparation of a nutritional
composition to reduce the uptake of high molecular weight substances, allergens and microorganisms through the intestinal wall,
more particularly to reduce transport of high molecular weight substances, allergens and microorganisms through the tight junctions
in the intestines, the rise in the viscosity of the nutritional composition caused by the polysaccharides being less than 20 mPa.s. The
nutritional compositions can be used to prevent or to treat allergy, allergic reactions, sepsis and inflammatory processes, such as
can arise under emotional and physical stress, ischaemia, reperfusion damage during and after operations, after radiation treatment
and/or chemotherapy of cancer patients and in the case of inflammatory diseases of the intestine, diarrhoea and allergies.

Nutritional compositions which contain non-digestible polysaccharides and use thereof to reduce transport through tight junctions

The present invention relates to nutritional compositions which contain a specific
5 class of non-digestible dextrans, hydrolysed (galacto)mannans and/or hydrolysed
(gluco)mannans. These compositions reduce the uptake of high molecular weight
substances, allergens and microorganisms through the intestinal wall. In particular, the
present invention relates to reduction of the free transport of such substances through the
tight junctions (TJ) of the intestines, without the transport of low molecular weight
10 substances, such as nutrients, via the intestinal epithelium being impeded. The
compositions can be used to prevent the increased permeability of the intestinal wall,
resulting from various causes, and the penetration of toxins, antigens and pathogenic
microorganisms present in the lumen which is caused as a result.

The structure and function of tight junctions is described, inter alia, in Ann. Rev.
15 Physiol. 60, 121-160 (1998) and in Ballard T.S. et al., Annu.Rev.Nutr., 1995, 15:35-55.
Tight junctions do not form a rigid barrier but play an important role in the diffusion
through the intestinal epithelium from lumen to bloodstream and vice versa.

The permeability of the tight junctions is highly regulated and can be disturbed by
illness and certain toxins in the lumen. Regulation takes place from the nervous system,
20 the hormonal system and the immune system. When the tight junctions open, substances
which have a high molecular weight, allergens and even microorganisms will pass through
the tight junctions. The translocation of substances having a high molecular weight can
under certain circumstances give rise to sensitisation of the immune system and result in
allergic reactions on subsequent exposure. Translocation of pathogenic microorganisms
25 imposes greater strain on the immune system and can make persons and animals ill, inter
alia in periods of lowered resistance. The same applies, for example, in the case of
bacterial toxins which have been able to pass through the epithelial layer and have been
able to reach the bloodstream.

The invention now relates to the use of one or more non-digestible polysaccharides
30 selected from the group consisting of dextrans having a molecular weight of 8 kD to
40,000 kD, hydrolysed (gluco)mannans having a molecular weight of 0.5 kD to 1,000 kD
and hydrolysed (galacto)mannans having a molecular weight of 0.5 kD to 1,000 kD to
reduce the uptake of high molecular weight substances, allergens and microorganisms

through the intestinal wall, with the proviso that the rise in the viscosity of the nutritional composition caused by the polysaccharides is less than 20 mPa.s.

More particularly, the invention relates to the use of the abovementioned compositions to reduce transport of high molecular weight substances, allergens and
5 microorganisms through the tight junctions in the intestines.

In addition to reducing the transport of harmful substances and microorganisms to a significant extent, a significant advantage of the present invention is that the normal transport of useful substances (nutrients) such as glucose, amino acids, dipeptides or trace elements is virtually maintained.

10 According to the invention non-digestible polysaccharides are understood to be polysaccharides which are not, or are barely, digested or converted by the human digestive enzymes under the conditions prevailing in the body. It should be pointed out that some of the non-digestible polysaccharides can be fermented by the microorganisms present in the intestines (colon, caecum and part of the ileum). Without wishing to be tied to any theory,
15 it is, however, expected that the effect of the polysaccharides on the paracellular transport does not take place via the fermentation products.

The degree to which the polysaccharides are digested can be established using the method as described in Minekus, M., Ph.D. Thesis, University of Utrecht, 1998, Development and validation of a dynamic model of the gastrointestinal tract, Section 2.
20 The polysaccharides according to the invention are less than 50% digestible and preferably less than 30% digestible.

Dextrans according to the invention are understood to be dextrans obtained via a (bio)synthetic route or naturally occurring dextrans. The molecular weight of such dextrans can be regulated by partial acid or enzymatic hydrolysis of the molecule followed
25 by repeated fractionation and precipitation with alcohol or ultrafiltration. These methods, which are known per se to those skilled in the art, must be carried out in such a way that the molecular weight of the dextrans falls within the cited range of 8 kD to 40,000 kD.

Dextrans having a molecular weight of 20 kD to 2,000 kD are preferably used.

The term (gluco)mannans is used to refer both to the mannans and the
30 glucomannans. The same applies in the case of the (galacto)mannans. Examples of galactomannans are guar gum, locust bean gum and tara gum. These (galacto)mannans and (gluco)mannans are used in the hydrolysed form. The molecular weights are between 0.5 kD and 1,000 kD.

Mixtures of dextrans, (galacto)mannans and (gluco)mannans can also be used.

The hydrolysed (galacto)mannans or (gluco)mannans according to the invention can be obtained by partial, but extensive, hydrolysis, for example with the aid of enzymes suitable for this purpose, by means of which substantial quantities of oligosaccharides
5 having a chain length of 3 to 5,600, preferably of 4 to 1,000, are produced.

The polysaccharides are preferably present in the preparation in an amount such that the concentration of these polysaccharides in the intestines is 0.1 to 20 g/l, preferably 0.5 to 10 g/l and preferentially 1 to 6 g/l. The minimum quantity of the active ingredient is determined in that a significant decrease in the transport through the tight junctions is
10 detected.

It is not necessary for the polysaccharides to be administered at that location where the paracellular transport is disturbed. The presence of the active component at a location somewhere in the intestines between the stomach and the affected location is sufficient.

Some of the polysaccharides used according to the invention have a viscosity-
15 increasing action which could prevent the absorption of nutritional components. The preparation must have a composition such that the normal transcellular transport is not impeded.

More particularly, the nutritional composition according to the invention has a viscosity of less than 100 mPa.s, preferably less than 40, but even more preferentially less
20 than 30 mPa.s. For the present invention it is important in particular that the polysaccharides, independently of the other constituents of the composition, have only a low viscosity-increasing effect. The viscosity-increasing effect of the active polysaccharides in the composition must be less than 20 and preferably less than 10 mPa.s and can be, for example, 3 mPa.s. Thus, the major proportion of the viscosity of the
25 product is caused by components other than the polysaccharides in the product.

The viscosity is determined by means of a Carri-med at a shear rate of 100 per second and at 20°C.

In the case of dry products the viscosity limits described above apply after reconstitution of the product.

30 In general, therefore, the type of polysaccharide (molecular weight) and the concentration thereof will be so chosen that an optimum combination of effectiveness and viscosity is obtained. Not only molecule size, but also degree of branching and degree of loading determine action, viscosity and/or fermentation behaviour.

The polysaccharides according to the invention prevent the free transport of high molecular weight substances, allergens and microorganisms through the tight junctions of the intestinal wall. In this context high molecular weight substances are understood to be the substances which under normal conditions are not able to pass through the tight junctions, or are able to do so only in minor amounts, and which can be assumed to have a toxic or allergenic action. These substances will in general have a molecular size of above 4,000 Dalton. Antigens, substances which activate the immune system, are in general peptides, which may or may not have been glycosided, frequently with a molecular weight of more than 10,000 Dalton. Allergens are antigens which produce an allergic reaction which usually is mediated via immunoglobulin E.

In this context microorganisms are understood to be in particular microorganisms which occur in the intestinal lumen. Thus, for example, under certain conditions overgrowth of microorganisms can take place in the small intestine, as a result of which tight junctions are to an increased extent exposed to these microorganisms.

According to another aspect of the invention foods or preparations are proposed which contain these non-digestible polysaccharides. These foods can be:

- complete foods;
- food supplements;
- health-promoting preparations; and
- tube feeds.

The compositions according to the invention can be used to prevent or to treat specific types of allergy, allergic reactions, sepsis and inflammatory processes, such as can arise under emotional and physical stress, ischaemia, reperfusion damage during and after operations, after radiation treatment and/or chemotherapy of cancer patients and in the case of inflammatory diseases of the intestine, diarrhoea and allergies.

The complete foods and food supplements described above can in particular be used in the treatment of, or to prevent, inflammatory diseases of the intestines, such as colitis ulcerosa, inflammatory bowel disease and Crohn's disease. Specific other constituents which can be incorporated in such foods and supplements are growth hormones, glutamine, n-3 LCPUFAs and the requisite contents of macro- and microingredients.

Furthermore, the foods according to the invention can be used before and after operations. Specifically, ischaemia and reperfusion damage to the intestine often occur during operations, as a result of which the tight junctions open. Introducing the

polysaccharides according to the invention into the intestines before and after the operation could prevent the uncontrolled paracellular transport. The administration of these polysaccharides can also be beneficial after chemotherapy.

5 In the case of diarrhoea a number of pathomorphological changes can also arise which are associated with an increased permeability of the tight junctions. These changes can arise with specific types of diarrhoea. The complete foods and food supplements according to the invention can be used to counteract the adverse consequences of this increased permeability.

10 The tight junctions can also open during stress, both of a physical nature (for example endurance sports) and of an emotional nature, as a result of which bacterial translocation takes place. Examples of emotional stress under which this takes place is the stress which arises during the transport of pigs to the slaughter house. Contamination of the meat can occur as a result. Another example is the stress that occurs when weaning piglets. The polysaccharides can be administered before the stress takes place, during
15 stress or after the stress has taken place.

With the aid of the polysaccharides according to the invention it is also possible to prepare preparations which are suitable for patients who have a food allergy, such as an allergy to cow's milk or to gluten. The increase in the permeability as a result of exposure to the allergen can be prevented. These preparations are of such composition that they do
20 not contain the said allergens.

The invention is explained on the basis of the following examples and with reference to the appended figures, in which

Figure 1 shows the Ussing chamber used in the examples;

Figure 2 shows the inhibition of the effects of caprate by dextran;

25 Figure 3 shows the inhibition by hydrolysed tara gum of the increased paracellular permeability caused by melitin;

Figure 4 shows the effect of dextrans on the HRP flow;

Figure 5 shows the effect of dextrans on the HRP flow in a pig under anaesthesia;

30 Figure 6 shows the effect of dextran on the increased permeability of the intestine of a microvillus inclusion patient.

Examples**I Examples of products**

Examples of compositions of various types of products in which the active component is dextran are given below.

- 5 The various types of product can be complete enteral foods, for use by the patient him or herself or for use as a tube feed. The product can be either in liquid form or in powder form, which is ready for use after dissolving. The active components can also be used as an ingredient in another food (for example bread) or in food supplements, such as a bar, a dairy product such as yoghurt, or a powder in the form of a sachet.

10

Example 1

Ready-to-feed, liquid, complete food for use before or after operations.

The composition is as follows per 100 ml of the product:

	Protein:	7.0 g
15	Fat:	4.0 g
	Carbohydrate:	21 g
	Dextran:	0.2 g

- 20 Minerals in a quantity of 1/15th of the recommended daily allowance (= RDA) can be added per 100 ml of the product. Trace elements and vitamins are added in somewhat larger amounts, i.e. 2/15 RDA. The product is of a composition such that 1,500 ml has to be consumed by the patient.

Example 2

- 25 Complete tube feed for persons suffering from inflammatory bowel disease. Per 100 ml the product contains:

Protein based on casein 7.0 g

Fat based on vegetable oils and 10% fish oil and 20% MCT; the linoleic acid content is 20% and the alpha-linolenic acid content 4.5%

- 30 Premixes containing the conventional forms of trace elements, vitamins and minerals Na, K, Ca, Mg, P, Zn, Fe, Mn, Cu, vit. B1, B2, niacin, A, D, K, B6, B12, pantothenic acid, folic acid.

Dextran: 0.6 g

Example 3

Food supplement for patients suffering from food intolerance or allergy.

Yoghurt based on soya milk. Per 100 ml the yoghurt contains:

Protein 4.0 g, fat 3.9 g, carbohydrates 12.3 g and 0.1 RDA of vitamins and trace elements.

Na = 80; K = 135; Cl = 125; Ca = 50; P = 50; Mg = 20 mg

Hydrolysed galactomannans 0.5 g

Example 4

10 Energy drink for athletes.

Per 100 ml the liquid contains

	Carbohydrate:	7.0 g
	Glucose:	0.2 g
	Fructose:	1.8 g
15	Lactose:	0.4 g
	Sucrose:	1.7 g
	Polysaccharides:	2.5 g
	Organic acids:	0.4 g
	Minerals:	
20	Na:	37 mg
	K:	17 mg
	Cl:	58 mg
	Ca:	8 mg
	Mg:	1 mg
25	Vitamin C:	15 mg
	Dextran:	0.1 g

Example 5

Premix for use in pig or piglet feed.

30 A/Premix consisting of 90% cornflour and 10% 150 kD dextran

B/Premix consisting of a suitable premix of vitamins, trace elements and minerals and 10% dextran.

Premix A or B, or mixtures thereof, can be used in the production of pig feeds. These can

be special feeds for use when pigs are transported, have to be rehoused in the sty or if they have a period of lowered resistance.

The premixes can also be used in a piglet feed for use after weaning, as an additive or instead of the premixes which are already known for use in piglet feed.

5

II Effect on transport via the tight junctions of the intestine

Use was made of a model set-up for determination of the effect of the polysaccharides used.

10 A test animal, such as a rat or guinea pig, was brought under narcosis. The stomach wall was then opened and a piece of the ileum tied off. The intestinal tissue was removed and stripped of layers of muscle. The preparation thus obtained was then stretched between two compartments through which oxygenated solutions flowed (Figure 1). The preparation was treated either with buffer (control or blank) or caprate in buffer in order to open the tight junctions (100% permeability) or with the combination of caprate and a certain
15 concentration of polysaccharide in buffer. As a measure of the permeability the transport of HRP (horseradish peroxidase) over the preparation was determined in accordance with known methods.

The results of this type of experiments are shown in Figures 2 to 5.

20 The in vitro effect of dextran (70 kD) on the increased HRP flow caused by caprate in a guinea pig intestinal epithelium is shown in Figure 2.

The in vitro effect of hydrolysed tara gum (900 D) on the HRP flow of Caco-2 cells under the influence of 2 μ M melitin is shown in Figure 3. It can be seen that the increased paracellular permeability caused by melitin is inhibited by tara gum.

25 The effect of various dextrans on the HRP flow of Caco-2 cells under the influence of 2 μ M melitin is shown in Figure 4. Phar in the figure stands for Pharmacosmos.

Figure 5 shows the effect of dextran on the increased HRP flow in the pig intestine caused by ischaemia. The figure relates to experiments with a pig under full narcosis, in which segments of the caudal section of the jejunum were taken. The effect of 5.6 g/l dextran (70 kD)(D) in the in situ ischaemia reperfusion model in pigs as a function of the
30 duration of ischaemia was determined in comparison with control (C) where no dextran was introduced into the lumen during ischaemia. A significant fall in the HRP flow under the influence of dextran was found compared with the control value.

Suction biopsies were taken from the duodenum of a child suffering from

microvillus inclusion disease (MVID). In the Ussing chamber these preparations displayed a four-fold increase in permeability to HRP compared with the normal value. After adding 70 kD dextrans to the luminal compartment of the Ussing chamber to give a concentration of 4.2 g/l the permeability was reduced to the normal level. No further HRP could be
5 detected in the paracellular spaces or tight junctions by means of electron microscopy. A corresponding result was obtained with dextrans having a molecular weight of 150 kD.

Figure 6 shows the result of this experiment with dextrans having a molecular weight of 70 kD. After 120 minutes a clear difference is detectable in the permeability with and without the addition of dextrans.

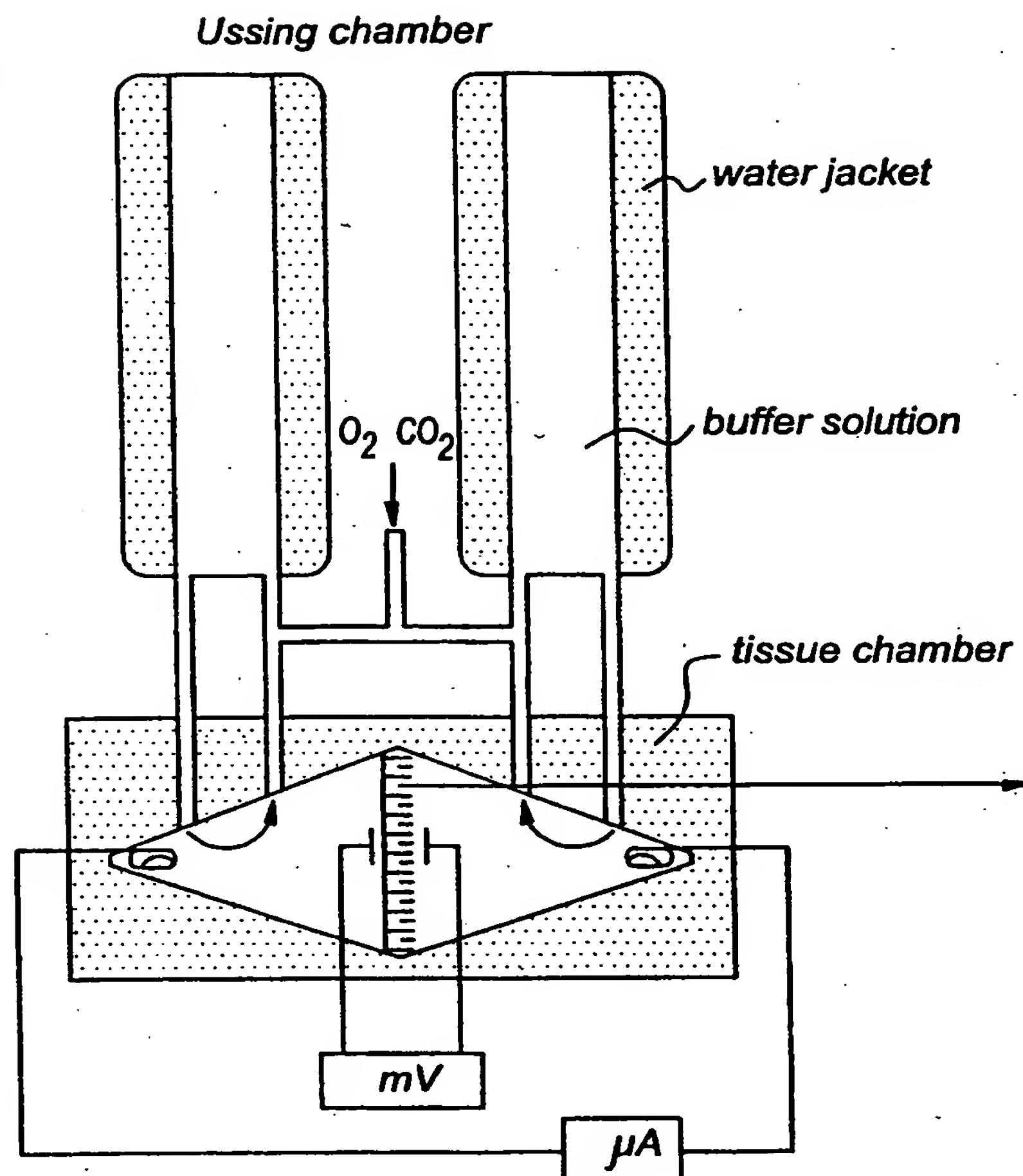
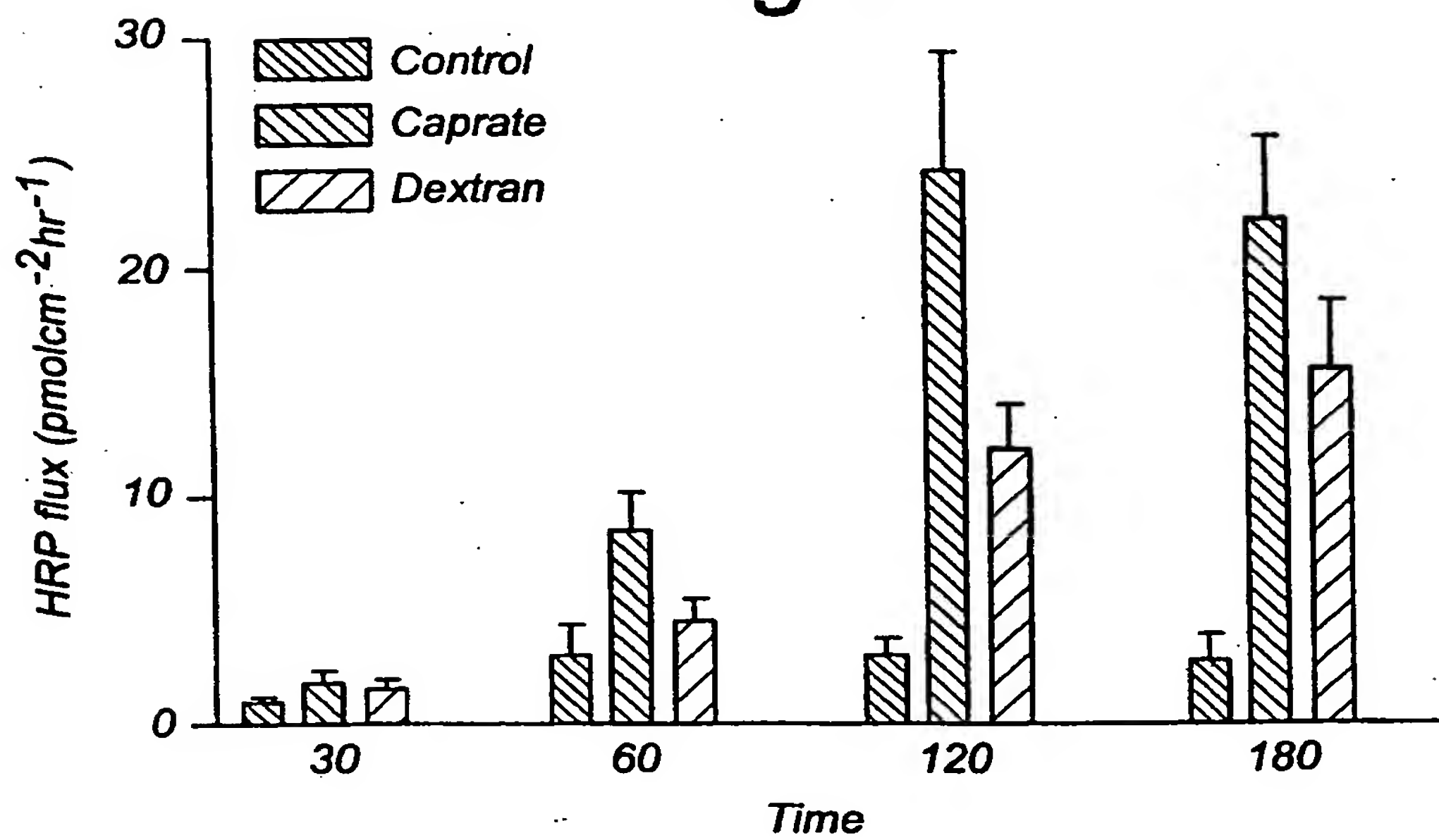
Claims

1. Use of one or more non-digestible polysaccharides selected from the group consisting of dextrans having a molecular weight of 8 kD to 40,000 kD, hydrolysed
5 (gluco)mannans having a molecular weight of 0.5 kD to 1,000 kD and hydrolysed (galacto)mannans having a molecular weight of 0.5 kD to 1,000 kD for the preparation of a nutritional composition to reduce the uptake of high molecular weight substances, allergens and microorganisms through the intestinal wall, with the proviso that the rise in the viscosity of the nutritional composition caused by the polysaccharides is less than
10 20 mPa.s.
2. Use according to Claim 1, wherein the polysaccharides are selected from dextrans having a molecular weight of 20 kD to 2000 kD.
- 15 3. Use according to one of the preceding claims, wherein the polysaccharides are contained in the composition in an amount such that the concentration of these polysaccharides in the intestine is 0.1 to 20 g/l, preferably 0.5 to 10 g/l and preferentially 1 to 6 g/l.
- 20 4. Use according to one of the preceding claims, wherein the nutritional composition is in the form of a complete food.
5. Use according to one of Claims 1 to 3, wherein the nutritional composition is in the form of a food supplement.
- 25 6. Use according to one of the preceding claims to reduce transport of high molecular weight substances, allergens and microorganisms through the tight junctions in the intestines.
- 30 7. Use according to one of the preceding claims, to prevent or to treat allergy, allergic reactions, sepsis and inflammatory processes, such as can arise under emotional and physical stress, ischaemia, reperfusion damage during and after operations, after radiation treatment and/or chemotherapy of cancer patients and in the case of inflammatory diseases

of the intestine, diarrhoea and allergies.

8. Nutritional composition which contains dextrans having a molecular weight of 8 kD to 40,000 kD, with the proviso that the rise in the viscosity of the nutritional composition
5 caused by the dextrans is less than 20 mPa.s.

1/3

Fig 1**Fig 2**

2/3

Fig 3

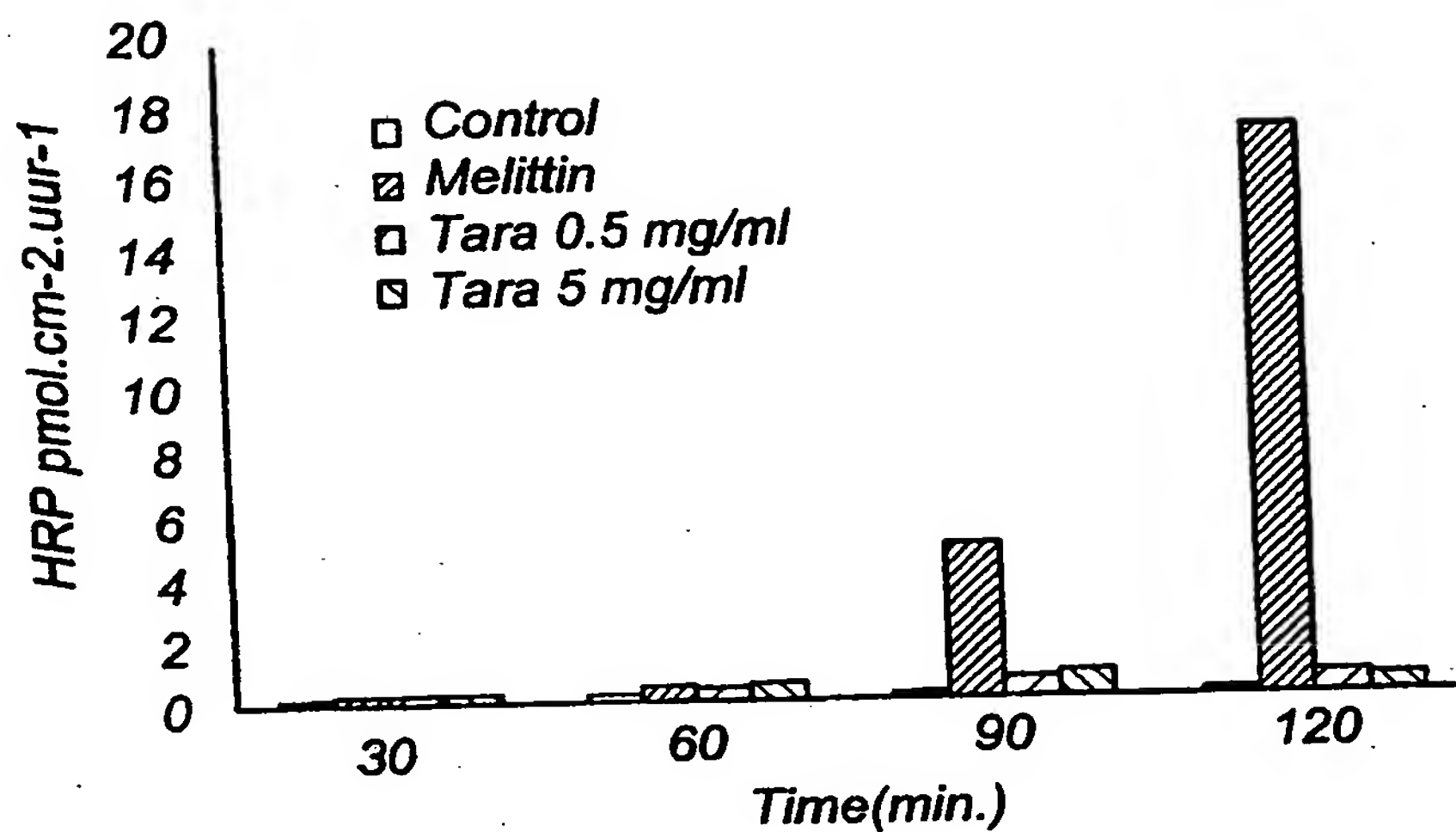


Fig 4

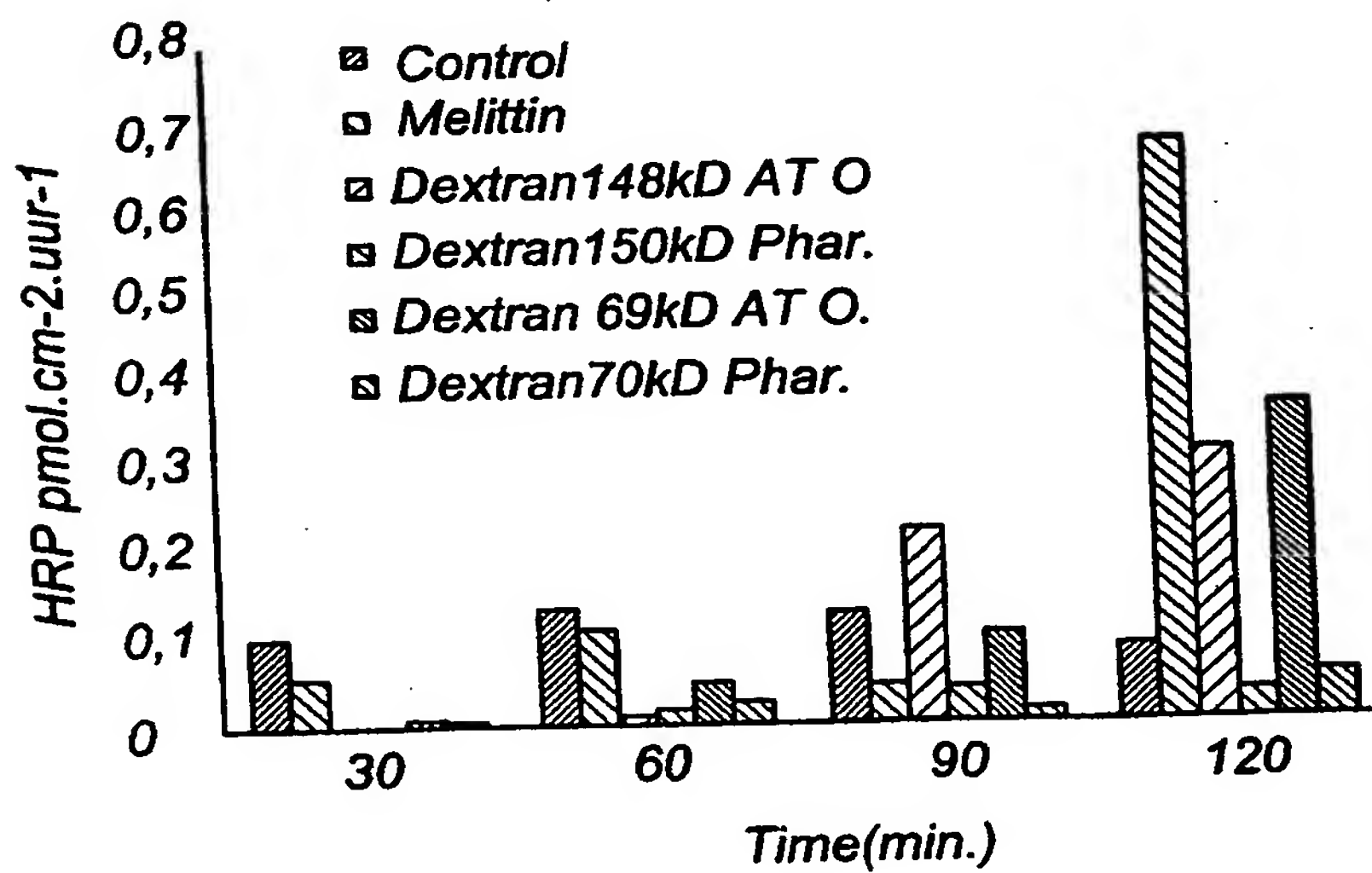
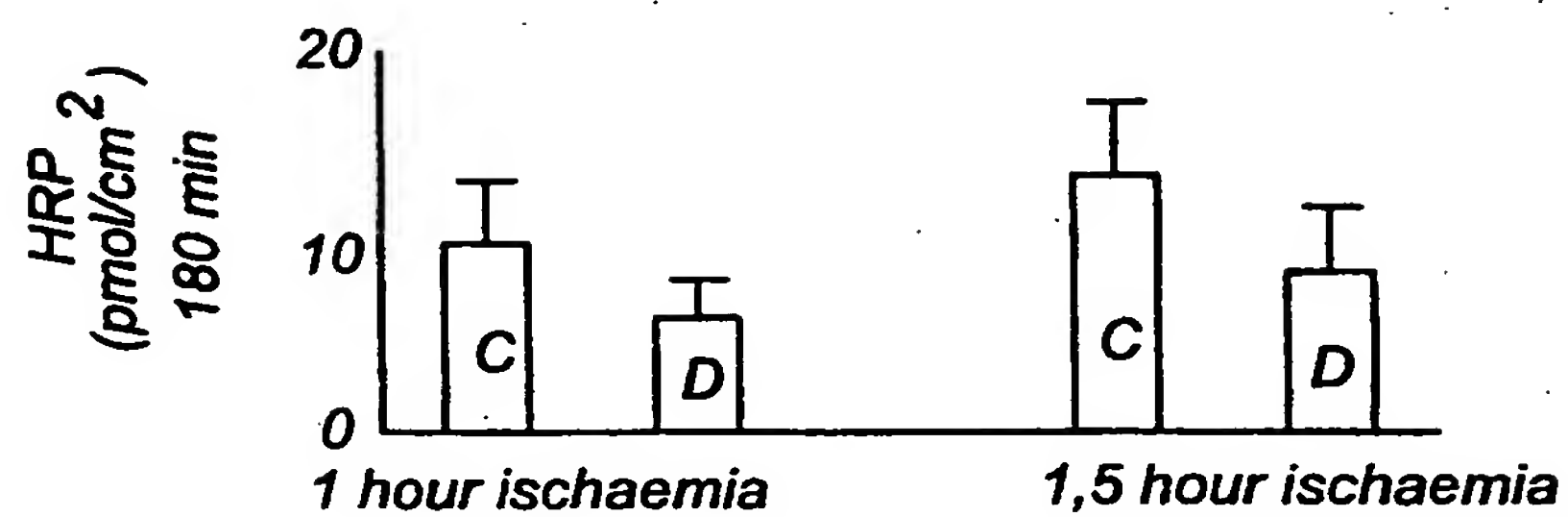
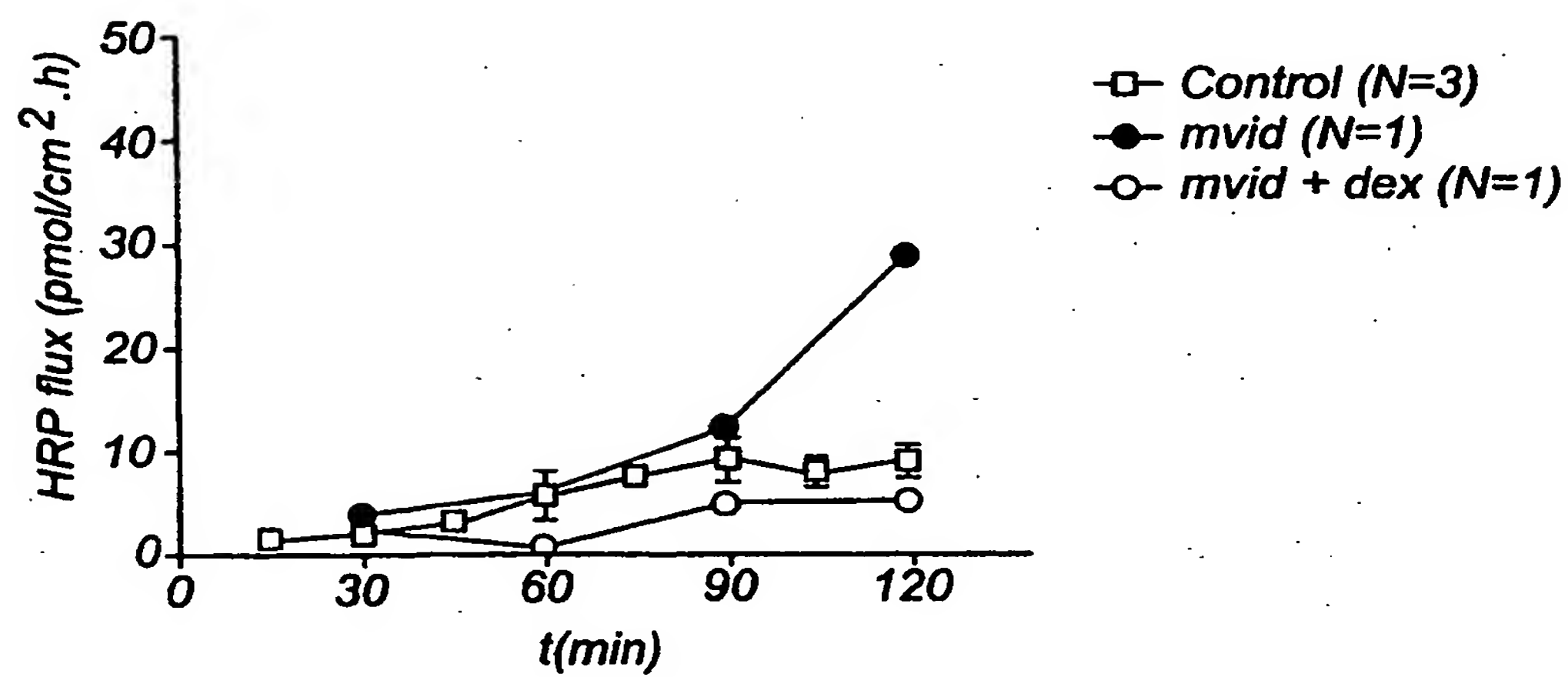


Fig 5*Fig 6*

INTERNATIONAL SEARCH REPORT

Int'l Application No
PCT/NL 00/00697

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A23L1/054 A23L1/0528 A23L1/0526 A61K31/715 A61P3/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A23L A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, PAJ, EPO-Internal, FSTA, BIOSIS, CHEM ABS Data, EMBASE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 153 013 A (FISONS PLC) 28 August 1985 (1985-08-28) page 2, line 22 -page 4, line 14 page 6, line 1 - line 13 page 7, line 1 - line 22	1-8
X	US 5 260 279 A (GREENBERG NORMAN A) 9 November 1993 (1993-11-09) column 2, line 20 - line 66	1-7
X	EP 0 385 598 A (MORINAGA MILK INDUSTRY CO LTD) 5 September 1990 (1990-09-05) page 2 -page 3, line 20 table 2	1-7
E	WO 00 57717 A (SAMOGGIA SIMONE ;LESEPIDADO S R L (IT)) 5 October 2000 (2000-10-05) the whole document	1-8

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *G* document member of the same patent family

Date of the actual completion of the international search

23 January 2001

Date of mailing of the international search report

01/02/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Vuillamy, V

INTERNATIONAL SEARCH REPORT

Information on patent family members

In International Application No

PCT/NL 00/00697

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0153013	A	28-08-1985	AT 56148 T	15-09-1990
			AU 571641 B	21-04-1988
			AU 3819385 A	08-08-1985
			DE 3579471 D	11-10-1990
			DK 39685 A	02-08-1985
			JP 60188403 A	25-09-1985
US 5260279	A	09-11-1993	AT 120930 T	15-04-1995
			AU 651626 B	28-07-1994
			AU 8603491 A	30-04-1992
			CA 2053933 A	25-04-1992
			DE 69108846 D	18-05-1995
			DE 69108846 T	12-10-1995
			DK 483070 T	17-07-1995
			EP 0483070 A	29-04-1992
			ES 2073154 T	01-08-1995
			JP 4282316 A	07-10-1992
EP 0385598	A	05-09-1990	JP 2222659 A	05-09-1990
			JP 2639726 B	13-08-1997
			DE 69002482 D	09-09-1993
			DE 69002482 T	10-02-1994
			US 4971814 A	20-11-1990
WO 0057717	A	05-10-2000	NONE	

THIS PAGE BLANK (USPTO)